requested. Attached hereto is a marked-up version of the changes made to the claims by the current amendment. The attachment is captioned "Version With Markings To Show Changes Made."

#### Rejection In View Of Priegnitz

Claims 15-23 have been rejected under section 103 as being unpatentable over U.S. Patent No. 6,162,267 to Priegnitz et al. U.S. Patent No. 6,162,267 was filed on Dec. 11, 1998.

The Attached to this Amendment is a 131 Declaration (actually there are two identical 131 Declarations, one is signed by Dr. Yong Wang and the other signed by Drs. Tonkovich and VanderWiel). As established by the facts set forth in the attached 131 Declaration, applicants possessed the claimed invention on or before the effective date of the cited reference (Priegnitz). Alternatively, the attached 131 Declaration shows as much of the claimed invention as is shown in the cited reference (Priegnitz) and therefore the Priegnitz reference is eliminated as prior art. See MPEP 715.02 citing *In re Stryker* 168 USPQ 372 (CCPA 1971) and *In re Wakefield*, 164 USPQ 636 (CCPA 1970). More specifically, the evidence in the 131 Declaration shows reduction to practice of: a steam reforming catalyst structure comprising a spinel support and a steam reforming catalyst (Rh or Ni). The catalyst structure was subsequently tested and found to possess the properties recited in claim 15 and described in the Example.

Thus, the 131 Declaration establishes that applicants possessed the claimed invention on or before Dec. 11, 1998 (or, alternatively, on or before Dec. 11, 1998, applicants possessed at least as much of the claimed invention as the Priegnitz reference). Accordingly, the Priegnitz reference has

been eliminated as prior art, and, withdrawal of the rejection in view of Priegnitz is respectfully

requested.

Additionally, Priegnitz does not teach or suggest some of the claimed features (nor are they

inherent). For example, Priegnitz does not teach or suggest a catalyst structure having a magnesia

passivation layer as recited in claim 20, nor a catalyst made by impregnating with magnesia as

recited in claim 19 (such as by impregnating an alumina support with a solution of magnesium

nitrate). Thus, the claimed invention is additionally patentable based on this ground.

Conclusion

If the Examiner has any questions or would like to speak to Applicants' representative, the

Examiner is encouraged to call Applicants' attorney at the number provided below.

Respectfully submitted,

Date: 11 Feb. 2003

send correspondence to:

Frank S. Rosenberg

18 Echo Hill Lane

Moraga, CA 94556

fax. no. (925) 376-8429

By: Front S. Roebley

Frank S. Rosenberg Registration No. 37,068

tel: (925) 376-8416

-5-

## VERSION WITH MARKINGS TO SHOW CHANGES MADE

#### IN THE SPECIFICATION

The paragraph beginning at page 1, line 9, after "RELATED APPLICATIONS", has been replaced with:

This is a divisional of U.S. Patent Application Ser. No. 09/375,615, now U.S. Patent No.

[\_\_\_\_\_\_,] <u>6,284,217</u>, filed August 17, 1999. --

#### IN THE CLAIMS

Claim 15 has been amended as follows:

- 15. (Amended) A steam reforming catalyst structure comprising:
- a support comprising a spinel support; and

a steam reforming catalyst selected from the group consisting of rhodium, iridium, nickel, palladium, platinum, carbide of group [IVb] <u>VIb</u> and combinations thereof;

wherein the catalyst structure has stability such that, when tested in a packed bed at 900°C, with a feedstream consisting essentially of methane and steam at a 1:1 ratio of methane to steam, at a constant pressure and a contact time such that there is a hydrocarbon conversion of at least 50%, and measuring the CO selectivity, wherein between 26 hours time-on-stream and about 40 hours time-on-stream, the CO selectivity remains essentially unchanged and the hydrocarbon absolute conversion changes less than about 5%.



**PATENT** 



IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re Application of:

Wang et al.

Application No. 09/930,040

Filed: Aug. 14, 2001

For: METHOD AND CATALYST

STRUCTURE FOR STEAM

REFORMING OF A HYDROCARBON Art Unit: 1764

Examiner: B. Yildirim

Atty Docket: B-1482-DIV



# **DECLARATION PURSUANT TO 37 CFR § 1.131**

- 1. The attached document in which the first line reads "POx/SR" is a copy of a lab notebook page that was written on or before 11 December 1998. The entry "Yong: Ni or Rh (oxide) on MgO/Al<sub>2</sub>O<sub>3</sub> (spinel)" is a note from a meeting showing that, on or before 11 December 1998, Yong Wang suggested preparing a steam reforming catalyst comprising Ni or Rh (oxide) on a MgO/Al<sub>2</sub>O<sub>3</sub> spinel support.
- 2. The attached document in which the first line begins "Precoat of" is a copy of a lab notebook page that was written on or before 11 December 1998. This page shows that, on or before 11

December 1998, a MgO/Al<sub>2</sub>O<sub>3</sub> spinel support was made by impregnating alumina with a magnesium nitrate solution.

- 3. The attached document in which the first line begins "Rhodium onto" is a copy of a lab notebook page that was written on or before 11 December 1998. This page shows that, on or before 11 December 1998, we prepared a rhodium oxide on MgO/Al<sub>2</sub>O<sub>3</sub> spinel catalyst structure. The support referred to on this page is the MgO/Al<sub>2</sub>O<sub>3</sub> spinel support described in paragraph 2 above.
- 4. Subsequent testing of the catalyst described in paragraph 3 demonstrated the time-on-stream performance shown in Fig. 2 of the patent application.
- 5. The attached document in which the first line begins "NiO of" is a copy of a lab notebook page that was written on or before 11 December 1998. This page shows that, on or before 11 December 1998, we prepared a nickel oxide on MgO/Al<sub>2</sub>O<sub>3</sub> spinel catalyst structure. The support referred to on this page is the MgO/Al<sub>2</sub>O<sub>3</sub> spinel support described in paragraph 2 above.
- 6. All of the attached documents have been copied without change except that dates have been blanked out.

7. I declare that all of the above statements made of my own knowledge are true and all statements made on information and belief are believed to be true. I understand that willful false statements and the like are punishable by fine or imprisonment, or both (18 U.S.C. §1001), and may jeopardize the validity of the application or any patent issuing thereon.

Date: 2/10/2003	By: Yong Wang
Date:	By:
Date:	By:Anna Lee Y. Tonkovich

- POX (SR ( CONTO)
11 C) The second of the second
"SIGNIFICANT HOO FORMATION (H. SELECTIVITY ??)  WEXT STEP: ALKALL PROMOTION
NEXT STEP: ALKALI PROMOTION
-> LEE: POX MAYBE OFF THE TABLE' SOON -> FOCUS ON SR A
" CAN ADD MORE OZ TO INCR. SEECTIVITY, BUT CONV. WILL DECK
- COZ FAVORED AT 1+16HER CHA
-> YONG: Ni or Rh (oxide) ON MgO/AlzO3 (spinel)
-SR: (mid-low Rh loadings done) Rh/a-A/203 - 3= 420 @ 25 ms
" lower loading performs better (2.8% Rh)
" both have very low long. < 25% - Select. < 70%
" J-Alf3 RESULTS FROM EARLIER! PERF. MUCH BETTER (~10% Conv.) (Selec).
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- 5% Co, Fe, Ru / j-AlzO3 FOR MDS REMDY TOMORROW
- NEED SUPPORTS - YOUG! Strem, Aldrich? - OKY ALERS ASSESSMENT
entity.
- ERIC: FOAMS FOR YUFEI ARE READY - CERAMIC ADHESIONS TECHNIQUES
- SINCLE POX/SR REACTORS: ADAPT. TO USE SAME 1/2" ANNULAN TUL
- check Sizing / flow options
· CAT. SELECTION
CH <sub>4</sub> $\rightarrow$ SYNGAS $\rightarrow$ MeOH PCC SR? POX?
PCC SR? >> F75
YOx!
- <i>SR</i>
" literatue! precrows metals - Rh best
supports - all - Alog & Zroz & MgO best
promoters - alkali oxides
- MeOH
" thermo: low T high P
" System: low P, high ed. activity (oney lower T)
"commercial: Cu-Zn/AloG3 (ICI - Catalog) 30 ATM
-@ 120 msec.: 2.5 g MaOH/g. cat. h (~100% selectivity)
"literature: Rh, Pd - selective but not as activity  Project No
Entered By Date
Disclosed To and Understood By
Signed 1 Date
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	49,1663 51,1702g 2,0039g	51,2980 MgO 0 = 0,1278g
	(94) total w = 2.0039s total w = 2.1318 g want 0.1279 g of MgO using	Must not con Must not con Twith any or fro
	Want 0.1279 g of MgO using	Mg(NO3)2. 6 H20 Amo
	$\frac{M_{9}O}{M_{5}(N_{0})_{2}\cdot 6H_{2}O} = \frac{40.3}{256.4} = \frac{0.125}{M_{5}N_{1}}$	thate => 0.8137g
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-73	> Used 0.8158g Mg Nitrate and 4.63, (easily west into	nl water added solution
	Into over and used same temp program	70: Seg 5 Seg 6 Ramp to 900° Dwell at 3.0°/min 120 min
. /	6.0 % MgO Loading  1.88 5 of - 70 + 100 mesh pre con  ( ~ 0.75 g < 100 mesh)	ated support
	Check for incipient wettness with water	r issing -100 me
; ; ;	$\sim 0.52$ galfor. 2365 of support $\approx$	2.20 ml/s

Date of Work

7

- 1

Date\_

Project No.

Entered By\_

Disclosed To and Understood By

	Rhodium onto MgO Precoated J-AlzO3. 11 13% as RhO2 (for Mother to Sym Go)
13699-0	Dish Wt. catisapport Preconted Support from pg 7
,	37.0331g 37.6992g = 0.6661 g of 6% MgO pre conted Support
	(187) total wt = 0.6661 g total wt = 0.7656g => 0.0995 g RhOz
	$\frac{RhQ}{134.9} = \frac{Rh}{RhQ} = \frac{x}{.09955} = \frac{102.9}{134.9} \Rightarrow .0759 gRh$ needed
	Using Eugellard's 10.37 % Rh solution: 10.37% solu-= .0759 g Rh
	Sol'n = .7321g $0.7376 g Sol'n + Water = 1.3155$
<i>ا</i> ا	(alcined at 500° for 180 min (pg 9 prymm)
	Out of oven: 37.8060 => 0,1068 g RhO2 with 0.7729 g
	Out oven: 37.8060 => 0,1068 g RhOz with 0,7729 g Total 0.1068 PD/.7729 g RhO. Cotalyst = 13.82 % RhOz

Disclosed To and Understood By

-;

**PATENT** 

### IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re Application of:

Wang et al.

Application No. 09/930,040

Art Unit: 1764

Filed: Aug. 14, 2001

Examiner: B. Yildirim

For: METHOD AND CATALYST

STRUCTURE FOR STEAM

REFORMING OF A

HYDROCARBON

Atty Docket: B-1482-DIV

## **DECLARATION PURSUANT TO 37 CFR § 1.131**

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7. I declare that all of the above statements made of my own knowledge are true and all statements made on information and belief are believed to be true. I understand that willful false statements and the like are punishable by fine or imprisonment, or both (18 U.S.C. §1001), and may jeopardize the validity of the application or any patent issuing thereon.

Date:	

By: Yong Wang

Date: Feb. 11, 2003

David D. VondonWiel

Date: Fels 10, 2003

Anna Lee V Tonkovich

P.04

6147333301 925-5 8429 frank rosenberg Feb 10 22 10:16a + POX/SR ( ... conf'D) " Oz conv. ~ 100% - CIMTS CHE CONU. " SHENIFICANT HEO FORMATION (HE SITECTIVITY ??) ---- HEXT STEP: ALKALI PROMOTION \* -- LEE: POX MAYBE OFF THE TABLE SOON -- PEXUS ON SR AK ADD MORE O'S TO MACK SEECTIVITY , BUT COUL WILL DECK TOZ FAVORED AT HIGHER CH -> YORG: Ni on Rh (oxide) on MgO/ALO3 (spinel) \* -SR: (mid-low Rh loadings done) Rh/a-AlzOs - 3- 100 @ 25 mg higher loading performs better (2.8% Rh) " born have very low land. 125% - Selvet. 10% · J-ALO3 RESULTS FROM EARLIER! PERF. ANCH BETTER (-100% CONV.) (Selection) C 15 max " 25 mon 78 RA (Hz . WAYNE - CAT. PREP. - 5% Co, Fe, Ru / +- AlzO3 FOR MDS PEMPY TOMORDOW - NEED SUPPORTS - YOUR! Strem, Aldrich ? -- GIR AND AND THE PROPERTY OF EREC. - ERIL: FORMS FOR YOFET ARE READY - CEREBONC. ADDESSORS TECHMORES - SINCUE POR/SR RENCTORS: ADAPT. TO USE SAME 1/2" ANNULAN TUB .je - check sizing / Stow options · CAT. SELECTION CH+ + SYNGAS - MOH

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	(94) total wit = 2.00395 total with = 2.1318 g want 0.1279 g of MgO using Mg(NGS)2-CHiO Admirant $\frac{MgO}{Mg(NO_2)_2\cdot 6H_2O} = \frac{40.3}{256.4} = \frac{Q.(279.5)}{Mg(NO_2)_2\cdot 6H_2O} = 0.8137g$
	Mg(NO2):6H20 2564 Mg Nitrate => 0.81379
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ARCS.	Used 0.81585 Alg Althorte and 4: to 3 rol water added.
	Into oven and used same temp program to: Seg 5 Seg 6 Ramp to 9000 Dwell at 3:00/min 120 min
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0.7376 g sala + water = 1.3755

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Och oven: 37.8060 = 0,1068 g RhO. With 0.7729 5 0.1068 Bal/ .7729 5 RhO. Colify + 13.82 % RhOz

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8

AtiO of 13.75 onto Mg O Preconted & - Ala O3 (Method to Syn God) theing support made on previous page:

Dit & cal support

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(87.0%) totaled = 0.6577g = 0.0983g NiO wanted

=> 0.3826 5 A)(NO3)-LH20 model NO = 74.71 = 0.0983 NOOD UP = 290.81 - NIGOD LHO

2,20 1 x . 65775 = 1.45 ml

Ased > 0.3862 g in 1.44 ml Aco

Out of over, 500°: 38.48899 = .0909 g Ni O 0.7486 9 HA = 12.14% NiO

- Paper lists 7's in oxides

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